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- What is claimed is:
1. A method of ameliorating impaired glucose tolerance in a subject in need thereof, comprising: administering to the subject a pharmaceutical composition, comprising placental adherent stromal cells (ASC), thereby ameliorating impaired glucose tolerance.
 2. The method of claim 1, wherein said subject is at least 60 years of age at the onset of treatment.
 3. The method of claim 1, wherein said subject has a HbA1c value of at least 43.5 mmol/mol at the onset of treatment.
 4. The method of claim 1, wherein said subject has a body mass index of at least 27.5 kg/m² at the onset of treatment.
 5. A method of reducing systemic inflammation in a subject with impaired glucose tolerance in a subject in need thereof, comprising: administering to the subject a pharmaceutical composition, comprising placental adherent stromal cells (ASC), thereby reducing systemic inflammation in a subject with impaired glucose tolerance.
 - 6-8. (canceled)
 9. The method of claim 1, wherein said ASC have been incubated on a 3D culture substrate in a bioreactor.
 10. The method of claim 9, further comprising the subsequent step of harvesting said ASC by removing said ASC from said 3D culture apparatus.
 11. The method of claim 9, wherein said ASC have been incubated on a 2D adherent-cell culture substrate, prior to incubation in said 3D culture apparatus.
 12. (canceled)
 13. The method of claim 9, wherein said 3D culture substrate comprises a synthetic adherent material formed as a fibrous matrix, wherein said synthetic adherent material is selected from the group consisting of a polyester, a polypropylene, a polyalkylene, a polyfluorochloroethylene, a polyvinyl chloride, a polystyrene, a polysulfone, a cellulose acetate, a glass fiber, a ceramic particle, a poly-L-lactic acid, and an inert metal fiber.
 - 14-16. (canceled)
 17. The method of claim 1, wherein said administering comprises:
 - a. administering to the subject a first pharmaceutical composition, comprising allogeneic placental ASC from a first donor; and
 - b. administering to said subject, at least 7 days after step a), a second pharmaceutical composition comprising allogeneic placental ASC from a second donor, wherein said second donor differs from said first donor in at